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SHORT COMMUNICATION

Evaluation of Buffalograss Leaf Pubescence and Its Effect on Resistance to Mealybugs (Hemiptera: Pseudococcidae)

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Introduction

Considerable progress has been made in improving the turfgrass characteristics of buffalograss, *Buchloë dactyloides* (Nutt) Engelm, a native North American grass species with low maintenance requirements (Riordan *et al.*, 1993). Two mealybugs, *Tridiscus sporoboli* (Cockerell) and *Trionymus* sp. (Hemiptera: Pseudococcidae) have emerged as buffalograss pests Baxendale *et al.*, 1994).

Mealybugs have been associated with buffalograss stands throughout Nebraska (Baxendale *et al.*, 1994), as well as in Texas and Arizona. Unfortunately, the Pseudococcidae are poorly described and species identification requires extensive specimen preparation (Ferris, 1950, 1953). Buffalograss mealybugs have an oblong, pale purple-grey, membranous body, ranging in length from 0.2 to 3.0 mm. They are covered with cottony wax secretions (Baxendale *et al.*, 1994). The adult female is relatively immobile and is found inside or near the leaf sheath, or behind the leaf axils enclosing the female flower. Clusters of eggs are deposited within a filamentous waxy ovisac. First instars, or crawlers, migrate to new feeding sites and likely play a role in host selection (McKenzie, 1967). Mealybugs are often overlooked in the field because of their small size and hidden location on the plant (Baxendale *et al.*, 1994). Severe mealybug infestations result in a general decline of the buffalograss stand, which can be confused with drought or other stresses. Initially, the turf takes on a reddish-purple discoloration, followed by browning and thinning. A close examination will reveal the mealybug's white cottony secretions.

Possible strategies for managing mealybugs on buffalograss include pesticides, biological control, and use of resistant buffalograsses. Developing mealybug-resistant buffalograsses is of particular importance because this grass is used primarily as a low-input turfgrass species. Fortunately, several resistant buffalograsses have been identified (Johnson-Cicalese *et al.*, 1998). Understanding the mechanism of this resistance would be helpful for formulating optimal strategies for identifying and exploiting new sources of resistance. While considerable progress has been made in identifying germplasm resistant to insect pests, progress toward characterization of the mechanisms conferring the resistance has been limited.

Resistance mechanisms identified in other turfgrass species include: increased tolerance due to greater rhizome numbers, higher stored food reserves, improved turf density and plant vigor; antibiosis factors which reduce survival or oviposition of the insect, for example, endophyte-infection (*Acremonium* spp.) of the host plant; and antixenosis factors such as leaf size and width, and time of flowering that affect oviposition (Reinert, 1982; Quisenberry, 1990; Johnson-Cicalese *et al.*, 1989). In buffalograss, glabrous leaf surfaces are suspected of playing an important role in mealybug resistance (Johnson-Cicalese *et al.*, 1998). Two glabrous buffalograsses, '609' and 'Prairie', have been shown to be highly resistant to mealybugs, and significant positive correlations were found between leaf pubescence and mealybug infestation levels in several greenhouse screening trials (Johnson-Cicalese *et al.*, 1998).

Pubescence is widely recognized as a factor in plant resistance to insects. Norris and Kogan (1980) provided 55 examples of how pubescence affects the behavioral and physiological response of arthropods to plants. In a third of the cases cited, pubescence increased the susceptibility of the host plant by making it more suitable for oviposition, affecting feeding behavior, or providing protection from predators. Pubescence enhanced oviposition by several lepidopterous insects on a number of plant species (Lambert *et al.*, 1992; Navasero and Ramaswamy, 1991). Pubescent wheats (*Triticum* spp.) were more heavily infested by airborne wheat curl mites (*Eriophyes tulipae* Keifer) (Harvey and Martin, 1980). Among several

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wheatgrass species (*Agropyron* spp.), pubescence seemed to increase the level of resistance to the black grass bug (*Labops hesperius* Uhler) (Ling *et al.*, 1985).

A number of researchers have employed scanning electron microscopy (SEM) to characterize the leaf surface and pubescence in plant resistance (Ling *et al.*, 1985; Navasero and Ramaswamy, 1991; Dahlin *et al.*, 1992) and grass morphology studies (Sangster *et al.*, 1983; Chen and Fukuoka, 1991). SEM has also been used to study mealybug anatomy (Cox and Pearce, 1983). The objectives of this study were to use light and scanning electron microscopy (SEM) to disclose morphological differences between mealybug-resistant and susceptible buffalograsses, and to enhance our understanding of the underlying resistance mechanisms.

Materials and Methods

TRICHOME COUNTS AND PUBESCENCE RATINGS. Trichome counts were taken on 16 buffalograsses exhibiting a range of mealybug infestation levels and pubescence ratings (Johnson-Cicalese *et al.*, 1998). These counts were taken to document trichome densities differences among buffalograsses, and to confirm the accuracy of correlations between trichome numbers and pubescence ratings. Pubescence ratings, while subjective, were time-efficient, taking only a few seconds to evaluate each plant. The rating scale was 1–6, where 1 = no trichomes and 6 = very dense trichomes (approximately 30% of the leaf surface covered) (Johnson-Cicalese *et al.*, 1998).

Plants were maintained under uniform conditions in the greenhouse, under 400-watt HID lamps [photoperiod of 16:8 (L:D) h] and at maximum/minimum temperature settings of 27/21°C. A fully expanded leaf blade was removed from two different pots of each buffalograss and examined. Trichomes seen within an ocular grid at 30× magnification were counted. Depending on leaf width the area observed was approximately 4 mm². To determine if trichome density varied from one location on the leaf to another, both the abaxial and adaxial leaf surfaces of an area adjacent to the ligule and 20 mm from the ligule were examined.

Experimental design was a split-plot; the main unit treatments were 16 buffalograsses in a completely randomized design (two replications), and the subunit treatments were the four locations on the leaf surface. Data were subjected to analysis of variance, and when a significant *F* ratio occurred ($P < 0.05$), means were compared using the least significant difference test (LSD). Correlations between counts and ratings were also analyzed (SAS Institute, 2002).

SCANNING ELECTRON MICROSCOPY (SEM). SEM was used to examine buffalograss leaf pubescence in greater detail, and to evaluate its possible effect on susceptibility to mealybugs. Two mealybug-susceptible buffalograsses, 85-97 and 378, and two mealybug-resistant buffalograsses, 609 and *Prairie* plants were examined. Prior to examination, plants were maintained in the greenhouse under the same conditions as previously described. Because mealybugs are typically found on the adaxial surface at the junction of the leaf blade and sheath, this was the primary area examined. The second fully-expanded leaf was examined in all cases. To evaluate mealybug interactions with the plant, nymphs, adults, and mealybug-infested leaves were also examined.

Previous SEM studies involving grasses and mealybugs have utilized several methods of specimen preparation (Cox and Pearce, 1983; Sangster *et al.*, 1983; Chen and Fukuoka, 1991). In this study, two methods were used, fixed and fresh tissues. For fixed tissues, a 10 mm leaf segment was immersed in 3% glutaraldehyde in a phosphate buffer for 1 h, dehydrated in an ethanol series (20%, 50%, 75%, 95%, 100%), then critical point dried with CO₂ in a Sorvall Critical Point Drying System (Sorvall, Newton, CN 06470). Leaf segments were attached to stubs using double-sided adhesive and immediately coated with 300A Au or Au/Pd in a Denton Desk II cool sputter coater (Denton Vacuum Inc., 1259 North Church St., Moorestown, NJ 08057). Fresh tissues were removed from the plant, mounted directly on stubs and coated similarly to the fixed tissues. Fixed and fresh mealybug specimens were prepared in a similar manner. Specimens were examined under a Cambridge S-90 Stereoscan electron microscope (Leo Electronic Microscopy Inc., One Corporation Way, Peabody, MA 01960).

Results and Discussion

TRICHOME COUNTS AND PUBESCENCE RATINGS. Significant differences in trichome densities were found among the buffalograsses ($F = 15.94$; d.f. = 15, 16; $P < 0.0001$), with number of trichomes ranging from 0 to 11/mm² (Table 1). The buffalograsses 85-97 and 378 were highly pubescent. The majority of buffalograsses, however, were intermediate in trichome density, and a few buffalograsses, including 609, 'Buffalawn', 84-412, and *Prairie*, had almost no trichomes. There were significantly more

Table 1. Trichome numbers and pubescence ratings of 16 buffalograsses.

Selection	Mean Trichomes*		Pubescence
	Abaxial $\pm$ SEM [†]	Adaxial $\pm$ SEM	Ratings [‡]
<i>Buffalawn</i>	0.0 $\pm$ 0	0.1 $\pm$ 0.1	1.0
'609' [§]	0.0 $\pm$ 0	0.0 $\pm$ 0	1.1
84-412	0.0 $\pm$ 0	0.2 $\pm$ 0.2	1.0
<i>Prairie</i>	0.1 $\pm$ 0.1	0.2 $\pm$ 0.1	1.7
84-714	0.2 $\pm$ 0.1	4.1 $\pm$ 1.8	2.8
85-204	3.3 $\pm$ 0.2	3.6 $\pm$ 0.5	4.0
85-25-2	4.8 $\pm$ 1.5	2.9 $\pm$ 0.7	4.0
'315'	7.7 $\pm$ 1.0	5.4 $\pm$ 0.6	3.4
84-931	6.6 $\pm$ 1.1	4.4 $\pm$ 1.0	4.8
85-217	6.4 $\pm$ 0.6	5.1 $\pm$ 0.9	5.0
84-22-2	6.0 $\pm$ 0.3	5.5 $\pm$ 0.8	4.8
84-924	8.6 $\pm$ 1.5	5.9 $\pm$ 1.0	3.8
84-WS	8.8 $\pm$ 1.7	4.9 $\pm$ 1.1	4.5
85-33	7.4 $\pm$ 0.5	6.2 $\pm$ 0.5	4.8
'378' ^{**}	10.9 $\pm$ 1.4	5.2 $\pm$ 0.9	4.5
85-97	8.7 $\pm$ 1.1	7.7 $\pm$ 1.1	5.6
LSD(.05)	2.2	1.7	0.6
Mean	5.0	3.8	3.6

* Mean number (adjacent to and 20mm from ligule) of trichomes per mm² of leaf blade.

[†] Standard error of the mean.

[‡] Pubescence ratings of whole plants (1–6 scale, where 1 = none).

[§] Mealybug-resistant standard.

^{**} Mealybug-susceptible standard.

trichomes on abaxial surfaces of leaf blades than on adaxial surfaces ($F = 16.12$; d.f. = 1, 80; $P < 0.0001$), although the differences were relatively small (abaxial = 5.0/mm², adaxial = 3.8/mm²). On the adaxial side, significantly more trichomes were found adjacent to the ligule (4.4/mm²) as opposed to 20 mm from the ligule (3.3/mm²) ($F = 15.48$; d.f. = 1,17; $P < 0.0011$); whereas on the abaxial side, the difference was not significant (adjacent: 5.1/mm², 20 mm: 4.8/mm²) ($F = 0.99$; d.f. = 1,16; $P < 0.34$). These relatively small differences among the four locations suggest that trichomes are uniformly distributed over the leaf surface (data not shown for the four locations). A primary reason for taking trichome counts was to compare trichome densities with pubescence ratings. Highly significant positive correlations were found between pubescence ratings and trichome numbers in all four positions on the leaf (abaxial/adjacent: $r^2 = 0.79$, $P = 0.0002$; abaxial/20 mm: $r^2 = 0.85$, $P = 0.0001$; adaxial/adjacent: $r^2 = 0.90$, $P = 0.0001$; and adaxial/20 mm: $r^2 = 0.89$, $P = 0.0001$). These results indicate that a visual assessment of the whole plant, using a 1–6 rating scale, is an accurate and efficient method for assessing plant pubescence.

Leaf width measurements were also taken for the 16 buffalograsses (data not shown), and no significant differences were found ($F = 1.19$; d.f. = 14, 30; $P < 0.33$). The mean leaf width was 1.7 mm and the range was 1.2–2.1 mm. Since leaf blade width has been shown to affect resistance to insects in other grass species (Ahmad and Funk, 1982), it was useful to eliminate this variable as a possible resistance factor in buffalograss.

Of particular interest in these pubescence evaluations was our observation that most of the buffalograsses with glabrous leaves had a lower ploidy level, and exhibited poor winter hardiness in Nebraska field trials. Geographic distribution of buffalograss is believed to be related to its ploidy level, with diploid buffalograsses occurring only in Central Mexico and Texas, tetraploids occurring in Mexico, Texas and a narrow band along the eastern slope of the Rocky Mountains, and hexaploids primarily throughout the central and northern Great Plains (Huff *et al.*, 1993). The diploid cultivar, *Buffalawn* (Table 1) was glabrous. This diploid suffers severe winter injury in Nebraska (Riordan *et al.*, 1995). Both *Prairie* and 609 have a tetraploid chromosome number, are free of pubescence (Table 1) and lack winter hardiness. The ploidy level of the glabrous buffalograss, 84-412 (Table 1), is unknown but was originally collected in Dallas, TX. All of the other selections evaluated are pubescent and believed to be hexaploids. They are all relatively winter hardy.

Fig. 1. Buffalograss leaf surface. (A) Dense pubescence on adaxial leaf surface of mealybug-susceptible selection 85-97 (fresh tissue; scale bar = 0.1 mm). (B) Glabrous, adaxial leaf surface of resistant cultivar 609 (fresh tissues; scale bar = 0.1 mm). (C) Closeup of trichomes on abaxial leaf surface of 378 (fixed tissue; scale bar = 0.03 mm). (D) Waxy platelets and stomate on adaxial leaf surface of 609 (fixed tissue; scale bar = 0.005 mm). (E) Glandular trichomes in ligule area of 609 leaf (fixed tissue; scale bar = 0.005 mm). (F) Shrunken glandular trichomes and leaf tissue of 609 (fresh tissue; scale bar = 0.02 mm).

Fig. 2. Mealybug/buffalograss interaction. (A) Mealybug nymph on 85-97 leaf (fixed tissue; scale bar = 0.1 mm). (B) Surface of mealybug illustrating waxy secretions, simple setae, and wax-producing pore (fixed tissue; scale bar = 0.004 mm). (C) 85-97 leaf surface with large quantities of mealybug wax adhering to the trichomes (fixed tissue; scale bar = 0.1 mm). (D) Close-up of mealybug wax on leaf surface, note long hollow tubes and curled filaments (fresh tissue; scale bar = 0.004 mm).

Trichome density and resistance to mealybugs may be related to ploidy level or adaptation. Both tetraploid selections (609 and *Prairie*) were highly resistant to mealybugs and 84-412 was moderately resistant (Johnson-Cicalese *et al.*, 1998). The association between mealybug resistance and ploidy level is generally consistent with the understanding that higher ploidy levels offer an advantage for improved adaptation due to the dosage effects of multiple alleles (Paterson, 2005).

SCANNING ELECTRON MICROSCOPY. SEM examination revealed numerous simple trichomes on the adaxial surface of the mealybug susceptible buffalograsses, 85-97 (Fig. 1A) and 378, and almost no trichomes on the mealybug resistant buffalograsses, 609 (Fig. 1B) and *Prairie* (SEM images not shown for 378 and *Prairie*). Raised cells surrounded the base of each trichome and the trichome surface was smooth (Fig. 1C). At higher magnification, a dense covering of waxy platelets was observed covering the leaf surface (Fig. 1D). This wax may be a factor in the drought resistance exhibited by buffalograss. Glandular trichomes were found in the ligule area of both the resistant and susceptible buffalograsses (Fig. 1E and 1F). The significance of these trichomes on buffalograss is unknown. In other plant species, glandular trichomes exude sticky or toxic substances and can be important in plant resistance to insects (Norris and Kogan, 1980).

A comparison of tissue preparation methods can be observed in Fig. 1E and 1F. The glandular trichomes on the fresh tissue sample (Fig. 1F) are shrunk and shriveled compared to the trichomes on the critical point dried sample (Fig. 1E). This shrunk appearance was observed in several fresh tissue samples. Fresh tissue is simpler to prepare, but this approach can result in distortion of samples and possible misinterpretation of results.

Examination of *in situ* mealybugs with light and scanning electron microscopy (Fig. 2) provided valuable clues to how this insect interacts with trichomes and the leaf surface. Young nymphs (crawlers) fit

between the trichomes, so pubescence probably does not hinder movement or feeding (Fig. 2A). In fact, the trichomes may actually provide a foothold. The abundant waxy secretions produced by mealybugs may interact with leaf trichomes and thus provide a more suitable environment for feeding and oviposition (Fig. 2B–D). Close-ups of the mealybug's cuticle reveal curved filaments of secreted wax and a wax-producing pore (Fig. 2B). Cox and Pearce (1983) discussed the function of the various forms of mealybug wax (see Fig. 2D). They suggested the curled filaments serve to protect the mealybug from their own honeydew by coating the droplets with wax. The long hollow tubes and shorter curled filaments are constituents of the ovisac and male cocoon. In buffalograss, leaf trichomes may provide a framework which supports the waxy filamentous ovisac and provides protection for eggs and newly emerged nymphs. Fig. 2C shows how mealybug wax has glued the trichomes together.

These SEM evaluations and a review of the literature (Norris and Kogan, 1980) suggest that pubescence may facilitate oviposition and provide a foothold for early-instar mealybugs. It is also possible that airborne mealybug crawlers are captured by the leaf trichomes, as occurs with the wheat curl mite (Harvey and Martin, 1980). Mealybug nymphs are known to climb to the upper leaves and tips of plants where they become airborne and are carried to new hosts (Nwanze, 1978). Leaf pubescence may also help prevent the parasitoid wasps that are known to be associated with these mealybugs (Heng-Moss *et al.*, 1998, 1999) from successfully parasitizing mealybug nymphs. Finally, highly pubescent leaves may alter the micro-environment of the leaf and provide a more favorable environment for mealybug development.

This research provides the first evidence that pubescent leaves increase buffalograss susceptibility to mealybugs, and suggests new avenues for further investigation. Specifically, future studies should focus on improving our understanding of the interactions among trichome density, ploidy level, adaptation, and resistance to mealybugs.

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